UP AND UP FAMOTIDINE- famotidine tablet, film coated
Target Corporation

Target Corporation Famotidine Tablets, 20 mg Drug Facts

Active ingredient (in each tablet)

Famotidine 20 mg

Purpose

Acid reducer

Uses

- relieves heartburn associated with acid indigestion and sour stomach
- prevents heartburn associated with acid indigestion and sour stomach brought on by eating or drinking certain food and beverages

Warnings

Allergy alert: Do not use if you are allergic to famotidine or other acid reducers

Do not use

- if you have trouble or pain swallowing food, vomiting with blood, or bloody or black stools. These may be signs of a serious condition. See your doctor.
- with other acid reducers

Ask a doctor before use if you have

- had heartburn over 3 months. This may be a sign of a more serious condition.
- heartburn with **lightheadedness, sweating, or dizziness**
- chest pain or shoulder pain with shortness of breath; sweating; pain spreading to arms, neck or shoulders; or lightheadedness
- frequent **chest pain**
- frequent wheezing, particularly with heartburn
- unexplained weight loss
- nausea or vomiting
- stomach pain
- kidney disease

Ask a doctor or pharmacist before use if you are

taking a prescription drug. Acid reducers may interact with certain prescription drugs.

Stop use and ask a doctor if

- your heartburn continues or worsens
- you need to take this product for more than 14 days

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Directions

- adults and children 12 years and over:
- to **relieve** symptoms, swallow 1 tablet with a glass of water. Do not chew.
- to **prevent** symptoms, swallow 1 tablet with a glass of water at any time from **10 to 60minutes before** eating food or drinking beverages that cause heartburn
- do not use more than 2 tablets in 24 hours
- children under 12 years: ask a doctor

Other information

- read the directions and warnings before use
- keep the carton. It contains important information.
- store at 20°-25°C (68°-77°F)
- protect from moisture

Inactive ingredients

carnauba wax, colloidal silicon dioxide, croscarmellose sodium, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, talc, titanium dioxide

Questions?

Call 1-888-547-7400

Principal Display Panel

see new warnings

Compare to active ingredient in Maximum Strength Pepcid® AC

maximum strength

famotidine tablets, 20 mg

acid reducer

just one tablet prevents and relieves heartburn due to acid indigestion

ACTUAL SIZE

50 + 50

100 TABLETS

2 x 50 TABLETS, 100 TOTAL

ACTUAL SIZE

see new warnings

NDC 11673-061-78

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Maximum Strength Pepcid® AC*

maximum strength
famotidine tablets, 20 mg
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up&up™

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Drug Facts (continued)

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GLUTEN FREE

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Minneapolis, MN 55403
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PAPER BOX
PLASTIC BOTTLE
Empty & Discard Cap
how2recycle.info

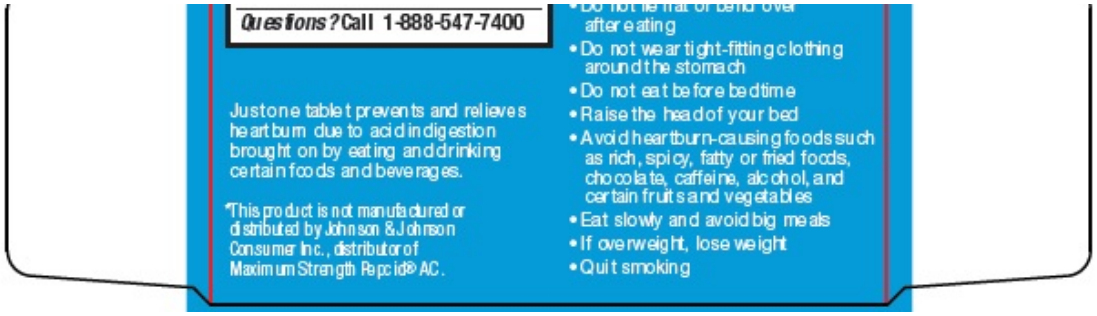
maximum strength
famotidine tablets, 20 mg
acid reducer

up&up™

DO NOT USE IF PRINTED FOIL UNDER CAP IS BROKEN OR MISSING

Drug Facts (continued)

Tips for Managing Heartburn



UP AND UP FAMOTIDINE

famotidine tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-061
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FAMOTIDINE (UNII: 5QZO15J2Z8) (FAMOTIDINE - UNII:5QZO15J2Z8)	FAMOTIDINE	20 mg

Inactive Ingredients

Ingredient Name	Strength
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	WHITE	**Score**	no score
Shape	ROUND	**Size**	8mm
Flavor		**Imprint Code**	L194
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:11673-061			

1	NDC:11673-061-71	1 in 1 CARTON	05/14/2012	
1		50 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:11673-061-02	25 in 1 CARTON	05/03/2012	
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:11673-061-72	1 in 1 CARTON	10/30/2013	10/10/2016
3		60 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:11673-061-78	2 in 1 CARTON	04/17/2020	
4		50 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:11673-061-82	1 in 1 CARTON	11/24/2021	
5		200 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077351	05/03/2012	

Labeler - Target Corporation (006961700)

Revised: 11/2021 Target Corporation